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This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c)

INVENTOR(S)

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Additional inventors are being named on the _____ separately numbered sheets attached hereto

TITLE OF THE INVENTION (200 characters max)

Odor-Absorbing Cellulosic Fibrous Substrates

Direct all correspondence to:

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ENCLOSED APPLICATION PARTS (check all that apply)

☒ Specification -- Number of Pages 10 CD(s) -- Number _____
☐ Drawing(s) -- Number of Sheets _____ ☒ Other (specify): Return postcard
☐ Application Data Sheet. See 37 CFR 1.76

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<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. <input type="checkbox"/> A check or money order is enclosed to cover the filing fees. <input type="checkbox"/> The Commissioner is hereby authorized to charge filing fees or credit any overpayment to Deposit Acct. No.: _____	FILING FEE AMOUNT (\$) \$ 160.00
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The invention was made by an agency of the United States Government or under a contract with an agency of the United States Government.

☒ No☐ Yes, the name of the U.S. Government agency & the contract number are: _____

Respectfully submitted,

Date: 11/05/2002

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ODOR-ABSORBING CELLULOSIC FIBROUS SUBSTRATES**Field of the invention:**

10 This invention is directed towards cellulosic fibers of fabric or other fibrous substrates coated with amines. These polymers impart durable anti-microbial activity, renewable control of certain odors, and the capacity to bind certain materials to the fabric surface.

Background of the invention:

15 Fabrics composed of only natural (e.g. cotton, wool, silk) or synthetic (e.g. polyester, nylon, acrylic) fibers are often lacking in desirable attributes. It is common in the textile industry to add a small weight component of various chemicals to the fabric to impart desired properties; these treatments are commonly referred to as "finishes". Such chemical enhancers include dyes, optical brighteners, softeners, water repellents, 20 water/oil repellents, insect repellents, anti-microbial and/or anti-fungal treatments, anti-static finishes, and hydrophilic finishes.

Durability is simultaneously a desired property and a significant challenge for any finish. Even molecules with only slight volatility will eventually evaporate; sunlight and air will slowly degrade others. Cleaning procedures such as laundering, dry- 25 cleaning, and shampooing are the most significant challenges to fabric finish durability. Many finishes are removed from fabrics after only a few cleanings.

Various approaches have been taken to provide durable finishes. One method is to deposit chemicals, typically polymers, that are not readily solubilized and washed away after being precipitated onto the fabric. Alternatively, the active ingredient of a 30 finish may be embedded in a laminant film that is applied to fabric; this procedure often allows for the slow release of the active ingredient into the surrounding fabric. However, the detergents and mechanical agitation of conventional cleaning procedures

often eventually remove the polymer or laminant film when it is merely deposited onto the fiber surface.

US Patent 6,187,856, issued to Incorvia et al, teaches the use of crosslinked resins, formed from polyamidoamines and polychlorohydrin crosslinkers, to form
5 durable films on fabrics. The resins of this patent are claimed to give durable anti-static properties to the fabric. Durability is defined in this patent as evidence of anti-static properties after dipping treated fabric into water heated at 80 °C for two twenty-minute intervals.

Anti-microbial finishes are highly desirable for many textile applications. They
10 may be employed on fabrics used in settings requiring antiseptic conditions, such as in hospitals. They may also be useful for fabrics worn or used in commercial food preparation, hospitality settings, and other areas where there is the significant potential of exposing people to infectious bacteria.

There are only a handful of classes of anti-microbial compounds. Durability is a
15 significant problem for them as most are small molecules that evaporate readily or can be washed away. Moreover, many anti-microbial compounds exhibit toxicity to humans. It would be desirable to invent a durable anti-microbial fabric finish that is innocuous to humans.

The ability to eliminate or significantly diminish malodorous axillary (body) odor
20 and foot odor is a desirable attribute for apparel fabrics. The chemical components of axillary odor are the waste by-products of certain bacteria that live off of the secretions from human sweat glands. These species of bacteria are called *lipophilic diptheroids*. Some three dozen molecules with potentially offensive odors have been identified in body odor (see, Preti, G. et al, *J. Chem. Ecology*, 1991, **17**, 1469; Preti, G. et al, *J.*
25 *Chem. Ecology*, 1992, **18**, 1039; Preti, G. et al, *J. Chem. Ecology*, 1996, **22**, 237; *Proc. Nat. Acad. Sci. USA*, 1996, **93**, 6626). All of them are organic acids and the main contributor to the odor has been identified as *trans*-3-methyl-2-hexenoic acid. The chemical components of foot odor have similar origin; they are waste products of the bacteria *brevidium epidermis*. These molecules are also organic acids, and the most
30 significant component is isovaleric acid (see, Kanda, F. et al, *Brit. J. of Dermatology*, 1990, **122**, 771). It would be desirable to have a durable finish that would eliminate or

significantly diminish malodorous body odor on fabrics. One approach is to include a bacteriocidal finish. However, these may not kill bacteria living on the skin and so odor may still be produced. Another method is to use a finish that absorbs the malodorous organic acids responsible for axillary and foot odor so that the volatile concentrations of the offensive organic acids are below the threshold of detectability. It would be greatly desirable to be able to recharge the absorptive capacity of such a finish by standard cleaning procedures.

US Patent 4,244,059, issued to Pflaumer, teaches the use of an odor-absorbent compound selected from alkali metal bicarbonates, alkali metal carbonates, water-soluble polyamines derived from ethylenimine, and mixtures thereof. The compound is deposited over the surface of air-permeable fabric composed of cellulosic fibers, to adsorb acidic and basic odorous molecules which are the major components of crotch odors. The patent makes no claims as to durability, nor does it make provisions to provide for durability of the polymer to the fabric during common cleaning processes such as laundering.

WO patent 97/34040, issued to Koizumi et al., teaches the use of polyamines as coatings for acrylic fibers to produce deodorizing fibers. In this patent, wet gel acrylic fibers containing acid groups are brought into contact with "an amino compound" with the stoichiometry adjusted so that there is an excess of amine groups. Electrostatic interactions between the amines and acid groups presumably are the source of durability. The fibers have been wet spun and not previously dried. After contacting the amine compound, the coated fiber is heated at between "100 and 180 °C under wet heat conditions." Fiber products constructed from these fibers are able to deodorize acidic odors.

SUMMARY OF THE INVENTION

This invention is directed towards durable finishes for cellulose-containing fibers and fibrous substrates. The active components of the finishes are hydroxyl-containing amines, and preferably trialkanol amines. When combined with a suitable crosslinker, the amines become attached to and crosslinked on the substrate fiber, and form a soft resinous coating that is durable to cleaning procedures.

This invention is further directed to the cellulosic fibers; yarns; woven, knitted or nonwoven fabrics and textiles; and finished goods (encompassed herein under the term "fibrous substrates") treated with the hydroxyl-containing amine coating of the invention.

5 The fibrous substrates treated with the finish described herein take on properties that are not found in the native fabric, including the ability to eliminate or greatly diminish the most offensive component of malodorous body odor, while reducing the yellowing of the substrates experienced with prior art amine treatments. Additionally, the treated cellulosic substrates remain hydrophilic and soft.

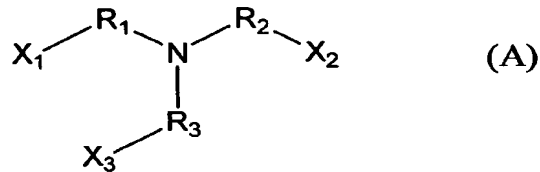
10 **DETAILED DESCRIPTION OF THE INVENTION**

As used herein and in the appended claims, "a" and "an" mean one or more, unless otherwise indicated.

15 The terms "durable" and "durability" as used herein describe a finished fibrous substrate in which the desired properties imparted to the substrate by the finish are observed after multiple launderings or dry cleanings.

20 The "cellulose-containing" or "cellulosic" fibrous substrates to be treated according to the present invention include any cellulosic fiber and any blend of fibers that contains a cellulosic, whether as a majority or a minority component. Cellulosic-based substrates include paper, cotton, rayon and other regenerated cellulose and cellulose-containing materials, linen, jute, ramie, industrial hemp, and the like. In a presently preferred embodiment, the cellulose-containing fiber or fibrous substrate is cotton.

25 The hydroxyl-containing amine may be a primary, secondary or tertiary amine, and may come from natural sources or from synthetic preparation. Tertiary amines are preferred because of their greatly reduced tendency to yellow compared to primary and secondary amines. Presently preferred embodiments of the invention include alkanol amines and preferably are selected from the trialkanol amines of Formula (A):



wherein, each of R_1 , R_2 and R_3 is independently selected from lower alkyl groups, unsubstituted or substituted with one or more hydroxyl groups; and each of X_1 , X_2 and X_3 is independently -OH or -H. Only one hydroxyl group per molecule is necessary for crosslinking of the molecule to the surface of the fibrous substrate. Having two or, preferably, all of X_1 , X_2 and X_3 being hydroxyl groups is desirable as it increases the likelihood of binding to the surface of the substrate and also allows crosslinking to other amines to improve the durability of the finish. The lower alkyl groups have from one to eight carbon atoms and may be straight-chained or branched.

Exemplary hydroxyl-containing amines useful in the present invention include, but are not limited to, triethanol amine; tris(hydroxymethyl)amino methane; 1-aza-3,3'-dioxabicyclo[3.3.0]octane-5-methanol; 1,3-bis[tris(hydroxymethyl)-methylamino]propane; and bis(2-hydroxyethyl)imino-tris(hydroxymethyl)methane.

The term "crosslinkers" as used herein describes molecules that contain two or more hydroxyl-reactive functional groups that form bonds with the hydroxyl groups on the hydroxyl-containing amine and on the cellulosic fibrous substrates. The crosslinkers bind the hydroxyl-containing amines together, as well as to bind the hydroxyl-containing amines directly to the fiber surface.

It is particularly desirable that the crosslinking reaction does not affect the basicity of the amines in the resulting coating. A catalyst may optionally be included to facilitate crosslinking. Hydroxyl-reactive functional groups include epoxides, halohydrins, oxiranes, carbonyl diimidazole, N,N'-disuccinimidyl carbonate or N-hydroxysuccinimidyl chloroformate, alkyl halogens, isocyanates, and N-methylol ureas. Preferred cross-linkers are diepoxides (Sigma-Aldrich corp.), N-methylol ureas, and blocked polyisocyanates such as Repearl MF (Mitsubishi Chemical Co.). Particularly preferred cross-linkers are the N-methylol ureas, such as dimethyloldihydroxyethyleneurea (DMDHEU) (PatCoRez P-53, BFGoodrich).

The finish that is applied to the fibrous substrate is a solution comprising at least a hydroxyl-containing amine, a cross-linker, and a volatile solvent. It is desirable that the amine and the cross-linker be soluble in the solvent. A particularly preferred solvent is water. The pad solution preferably contains hydroxyl-containing amine at
5 between about 0.01% and about 75% by weight, more preferably about 0.05% and about 50% by weight, and most preferably about 0.1% and about 20% by weight. The pad solution preferably contains a crosslinker at between about 0.001% and about 40% by weight, more preferably about 0.01% and about 30 % by weight, and most preferably about 0.05% and about 15% by weight. The finish solution may also include
10 other components as described below.

The reaction of the hydroxyl-containing amine with certain crosslinker functional groups, such as halohydrins, results in the formation of mineral acids that lower the pH of the finish and may slow the rate and decrease the extent of crosslinking. To control this deleterious effect, a buffering agent may be added to the finish solution. Buffering
15 agents are weak acids or bases that tend to hold solutions containing them within ± 1 pH point of the buffering agents' pK_a . One skilled in the art will appreciate that an optimal buffer solution consists of equimolar portions of the buffering agent and its corresponding conjugate acid or base, the latter often being formed by addition of a strong acid or base. Lists of buffering agents can be found in *Lange's Handbook of*
20 *Chemistry*, 14th edition, ed. J.A. Dean, McGraw-Hill, Inc., section 8, p.p. 103-112. If used, a buffering agent should be chosen so that the pK_a of the buffer lies within the optimal pH range of the reaction. This pH range is dependent on the identities of the reactive group of the hydroxyl-containing amine and of the crosslinker. The buffer must also be chosen so as to be unreactive with the crosslinker or the hydroxyl-containing
25 amine. The amount of buffering agent should be slightly more than equimolar to the theoretical total amount of acid generated by complete reaction of the crosslinker.

The finish solution may also include other additives. For example, amine groups are prone to oxidation, the by-products of which are often yellow-brown in color. While the finish of the present invention exhibits substantially less yellowing when tertiary
30 amines are used, it may be desirable to include the addition of an anti-oxidant to the finish solution to further minimize the extent of oxidation both in the stored state and

while the finish is being applied and cured on the fabric. A presently preferred anti-oxidant is phosphoric acid. Optical brighteners (such as, for example, the Leucophor products from Clariant Corp. or the Uvitex products from Ciba Corp.) may be included in the finish solution. Whitening agents, such as sodium tetraborate, may also be
5 included in the solution. The finish solution may also contain a wetting agent, such as WetAid NRW (BF Goodrich Corp.), to aid the equal spread of the finish over the fibers. Additional additives can be added to the solution as needed.

The finish can be applied to the cellulosic fibrous substrate by exposing the substrate to the finish solution by methods known in the art, such as soaking, spraying,
10 dipping, fluid-flow, and padding. The exposed fibrous substrate is then heated to remove the volatile solvent and to speed up the reaction of the hydroxyl groups on the substrate and in the hydroxyl-containing amine with the crosslinker. Alternatively, the cellulosic fibers or yarns may be exposed to the finish solution by soaking, spraying or dipping. After the finish is cured in place, the fibers or yarns may be woven or knit into
15 fabrics.

The finish solution may be applied to the fibrous substrate at any temperature above the freezing point and below the boiling point of the solvent. In the present embodiment, the application temperature is preferably between 5 and 90 °C, more preferably between 10 and 50 °C, and most preferably at room temperature. The
20 treated fabric should be cured at a temperature high enough to induce the crosslinking reaction in a very short time, preferably less than five minutes, more preferably a minute or less. In the present embodiment, the curing temperature is preferably between 100 and 200 °C, more preferably between 130 and 180 °C.

The present invention is further directed to the cellulosic fibrous substrates
25 treated with the finish described above. Substrates thus treated will possess properties not found in untreated substrates, while maintaining desirable properties such as a soft hand and hydrophilicity. These new properties include the ability to absorb malodorous organic acids via acid-base reactivity of the acids with the amine groups of the finish. The finishes of the invention are durable.

30 An embodiment of the present invention is the preparation of treated cellulosic fibrous substrates that absorb and deodorize organic acids, which gives such

substrates the ability to eliminate or greatly diminish offensive body odor. The odor-absorbing capacity of the fabric can be recharged when necessary by conventional laundering procedures. The molecular sources of offensive body odor are primarily the waste products of a species of bacteria named *lipophilic diphtheroids*. This species of bacteria lives on the skin surface of humans and primarily digests the secretions of the apocrine glands. The malodorous waste products of *lipophilic diphtheroids* are organic acids, with the most significant component being 3-methyl-2-hexenoic acid. Volatile organic acids are commonly considered to have highly offensive odors even in extremely low concentrations. The odor-absorptive capacity of the treated fibrous substrate stems from the basicity of the amine groups of the finish. Acids react with the free amine groups of the hydroxyl-containing amine to form non-volatile ionic complexes. The extent to which this ionic complexation occurs depends on the relative strength of the acid and base. In the case of the present invention, the reaction is biased towards formation of the ionic complex to such a degree that only between one acid molecule in ten thousand to one acid molecule in a million would be found in the non-ionized, potentially volatile form. Thus, as long as unreacted amine groups are available in the treated fabric, the concentration of volatilized organic acid around the treated fabric will be lowered to the point of being undetectable or scarcely detectable.

An advantage to the present invention over conventional odor-absorbing material such as activated carbon is the ability to recharge the odor-absorptive capacity of the fibrous substrate. As amines are weak bases, exposing the substrate to an aqueous solution with a pH at or above the pK_b of the base will deprotonate most of the amine complexes and result in separation of the amine-acid complexes. The conjugate base forms of the malodorous organic acids will be washed away in the laundry liquor, leaving behind free amine groups on the fiber surface. A pH of 10 is above the pK_b of most amines, and laundry detergent solutions such as Tide® typically have this pH or higher. Therefore, a conventional laundering procedure is sufficient to recharge the odor-absorptive capacity of the fabric.

The following examples are intended for illustrative purposes only. Those of skill in the art will recognize other embodiments, all of which are considered part of the present invention.

EXAMPLES

Example 1.

5 Samples (16x12 inch square) of untreated cotton twill fabric were immersed in
either a test solution (6.0 wt% triethanol amine, 10.0 wt% Patcorez P-53, and 0.25 wt%
Wet Aid NRW in water, final pH=4.0; Sample A) or in a control solution (water only,
final pH=4.0; Sample B), and padded at 30 psi. The samples were cured in a Mathis
oven set at 330°F (166°C) overall temperature with a 310°F (154°C) trigger
10 temperature for one minute. The whiteness of the resulting samples, compared to the
untreated fabric, was measured using a UV-Vis integrating sphere following AATCC
Test Method 110-2000. The samples were then home laundered ("HL") using 24 g. of
AATCC standard detergent in warm water on normal washer and dryer settings, after
which the whiteness index was again measured. The results are shown in Table 1
15 below.

TABLE 1

20

<u>Sample</u>	<u>Whiteness Index</u>					
	<u>0 HL</u>	<u>1 HL</u>	<u>5 HL</u>	<u>10 HL</u>	<u>15 HL</u>	<u>20 HL</u>
A	71.14	67.36	75.98	76.98	77.78	76.93
B	74.43	73.98	82.13	82.65	84.08	84.40

25

Hydrophilicity/hydrophobicity tests were run on the samples by measuring the
amount of time it takes for a drop of water to completely soak into the fabric. In all
cases with both the treated and control samples, the time was less than 2 seconds.

Finally, a smell test was run by placing solutions of various concentrations (in ppm) of butyric acid on the fabric samples and recording the lowest concentration of butyric acid that is noticeable on the sample by a panel of judges. These results are presented in Table 2, below.

5

TABLE 2. Smell Test Results

<u>Sample</u>	<u>Average minimum conc. (ppm)</u>					
	<u>0 HL</u>	<u>1 HL</u>	<u>5 HL</u>	<u>10 HL</u>	<u>15 HL</u>	<u>20 HL</u>
A	--	750	750	775	850	600
B	--	38	50	63	50	38

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